Abstract

SELF-MICROEMULSIFYING DRUG DELIVERY SYSTEMS OF A HIV PROTEASE INHIBITOR

The present invention relates to pharmaceutical formulations of (3R,3aS,6aR)-hexahydrofuro [2,3-b] furan-3-yl (1S,2R)-3-[[(4-aminophenyl) sulfonyl] (isobutyl) amino]-1-benzyl-2-hydroxypropylcarbamate, salts, esters, polymorphic and pseudopolymorphic forms thereof, which are self-microemulsifying drug delivery systems and comprise as carrier a lipophilic phase, one or more surfactants, a hydrophilic solvent and a nucleation inhibitor.